K001723

JUL 1 9 2000

510(k) Summary

Trade Name: **Cool-Tec Electrodes**

Sponsor: Seedling Enterprises, LLC
150 California Street
Newton, MA 02458
Registration No. not yet assigned

Device Generic Name: Electrosurgical electrode

Classification: According to Section 513 of the Federal Food, Drug, and Cosmetic Act, the device classification is Class II.

Predicate Devices: The proposed Cool-Tec Electrodes are substantially equivalent to several currently marketed electrosurgical electrodes which have been reviewed and cleared by FDA including the following:

EDGE Coated and Opti2 Electrodes (K962044); Valleylab, Inc., Boulder, CO
E-Z Clean Electrodes (K913473/K913281); MegaDyne Medical Products, Murray, UT

Product Description:
The Cool-Tec family of electrode probes are non-sticking electrosurgical devices that may be used with most commercially available electrosurgical generators. They eliminate sticking of tissue to the electrode by limiting the temperature of the metal electrode tip below the temperature at which tissue sticks to metal. This is done through a heat transfer means that takes the heat generated at the tissue, and transfers it to the handle for release to the air. These electrode probes may be used in both minimally invasive and open surgeries to replace currently available electrode probes.

Indications for Use:
Cool-Tec non-sticking electrodes are monopolar, electrosurgical devices intended for use in situations where monopolar electrosurgical cutting and coagulation are normally used.

Safety and Performance:
Substantial equivalence for this device was based on similarities in design and performance characteristics as well as performance testing. The materials, performance specifications and essential design characteristics of the Cool-Tec Electrodes are equivalent to those of the predicate devices.

Conclusion:
Based on the indications for use, technological characteristics, and comparison to predicate devices, the Cool-Tec Electrodes have been shown to be safe and effective for its intended use.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JUL 1 9 2000

Seedling Enterprises, LLC
c/o Ms. Pamela Papineau
Delphi Medical Device Consulting
5 Whitcomb Avenue
Ayer, Massachusetts 01432

Re: K001723
Trade Name: Cool-Tec Electrodes
Regulatory Class: II
Product Code: GEI
Dated: May 31, 2000
Received: June 6, 2000

Dear Ms. Papineau:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4595. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsmamain.html".

Sincerely yours,

Donna R. Bachman

Celia M. Witten, Ph.D., M.D.
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): K001723

Device Name: Cool-Tec Electrodes

Indications for Use:

Cool-Tec non-sticking electrodes are monopolar, electrosurgical devices intended for use in situations where monopolar cutting and coagulation are normally used.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Anne R. Rochner
vision Sign-Off)
sion of General Restorative Devices
(k) Number K001723

Prescription Use ✓ OR Over-the -Counter Use ____
(Per 21 CFR 801.109)